

INFORMATION DISCLOSURE STATEMENT	Atty. Docket No.: 232.00010120	Serial No.: 08/981,998
	Applicant(s): Pulst	
	Filing Date: May 11, 1998	Group: 1801 1655

U.S. PATENT DOCUMENTS

Examiner Initial	Document Number	Date	Name	Class	SubClass	Filing Date If Appropriate
	5,552,282	09/03/96	Caskey et al.	435	6	
	5,650,270	07/22/97	Giese et al.	435	6	
	5,650,277	07/22/97	Navot et al.	435	6	
	5,741,645	04/21/98	Orr et al.	435	6	

FOREIGN PATENT DOCUMENTS

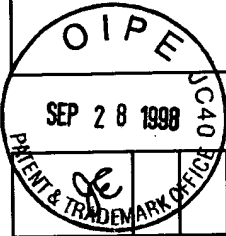
	Document Number	Date	Country	Class	SubClass	Translation	
						Yes	No
	WO 95/01437	01/12/95	PCT				
	WO 97/17445	05/15/97	PCT				X

OTHER DOCUMENTS (Including Authors, Title, Date, Pertinent Papers, etc.)

		Banfi, et al., "Identification and characterization of the gene causing type 1 spinocerebellar ataxia," <u>Nature Genetics</u> , 7, 513-519 (1994).
		Belal et al., "Clinical and genetic analysis of a Tunisian family with autosomal dominant cerebellar ataxia type 1 linked to the SCA2 locus," <u>Neurology</u> , 44, 1423-1426 (1994).
		Brook, "Retreat of the triplet repeat," <u>Nat. Genet.</u> , 3, 279-281 (1993).
		Brunner et al., "Brief Report: Reverse Mutation In Myotonic Dystrophy," <u>New Engl. J. Med.</u> , 328, 476-480 (1993).
		Filla et al., "Prevalence of hereditary ataxias and spastic paraplegias in Molise, a region of Italy," <u>J. Neurol.</u> , 239, 351-353 (1992).
		Gispert et al., "Chromosomal assignment of the second locus for autosomal dominant cerebellar ataxia (SCA2) to chromosome 12q23-24.1," <u>Nat. Genet.</u> , 4, 294-299 (1993).
		Imbert, "Cloning of the gene for spinocerebellar ataxia 2 reveals a locus with high sensitivity to expand CAG/glutamine repeats," <u>Nature Genetics</u> , 14, 285-291(1996).

EXAMINER 	Date Considered
*Examiner: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.	

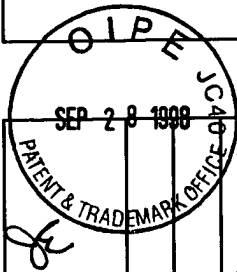
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		Ioannou et al., "A new bacteriophage P1-derived vector for the propagation of large human DNA fragments," <u>Nat. Genet.</u> , <u>6</u> , 84-89 (1994).
JE		Kawaguchi et al., "CAG expansions in a novel gene for Machado-Joseph disease at chromosome 14q32.1," <u>Nat. Genet.</u> , <u>8</u> , 221-227 (1994).
JE		Koide et al., "Unstable expansion of CAG repeat in hereditary dentatorubral-pallidoluysian atrophy (DRPLA)," <u>Nat. Genet.</u> , <u>6</u> , 9-13 (1994).
JE		Kremer, et al., "Mapping of DNA Instability at the Fragile X to a Trinucleotide Repeat Sequence p(CCG)n," <u>Science</u> , <u>252</u> , 1711-1714 (1991).
JE		Lopes-Cendes et al., "Confirmation of the SCA-2 Locus as an Alternative Locus for Dominantly Inherited Spinocerebellar Ataxias and Refinement of the Candidate Region," <u>Am. J. Hum. Genet.</u> , <u>54</u> , 774-781 (1994).
JE		MacDonald et al., "A Novel Gene Containing a Trinucleotide Repeat That Is Expanded and Unstable on Huntington's Disease Chromosomes," <u>Cell</u> , <u>72</u> , 971-983 (1993).
JE		Mahadevan, et al., "Myotonic Dystrophy Mutation: An Unstable CTG Repeat in the 3' Untranslated Region of the Gene," <u>Science</u> , <u>255</u> , 1253-1255 (1992).
JE		Mandel, "Questions of expansion," <u>Nat. Genet.</u> , <u>4</u> , 8-9 (1993).
JE		Nagafuchi et al., "Dentatorubral and pallidoluysian atrophy expansion of an unstable CAG trinucleotide on chromosome 12p," <u>Nat. Genet.</u> , <u>6</u> , 14-18 (1994).
JE		Orr et al., "Expansion of an unstable trinucleotide CAG repeat in spinocerebellar ataxia type 1," <u>Nat. Genet.</u> , <u>4</u> , 221-226 (1993).
JE		Polo et al., "Hereditary Ataxias and Paraplegias in Cantabria, Spain," <u>Brain</u> , <u>114</u> , 855-866 (1991).
JE		Pulst et al., "Anticipation in spinocerebellar ataxia type 2," <u>Nat. Genet.</u> , <u>5</u> , 8-10 (1993).
JE		Pulst et al., "Genetic and Physical Map of the Spinocerebellar Ataxia 2 (SCA2) Region on Human Chromosome 12," <u>Neurology</u> , <u>45</u> , A422 (1995).
JE		Pulst et al., "Moderate expansion of a normally biallelic trinucleotide repeat in spinocerebellar ataxia type 2," <u>Nature Genetics</u> , <u>14</u> , 269-276 (1996).
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EXAMINER <i>Jeanne Eneudo</i>	Date Considered <i>Jan 31, 2000</i>
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JE		Rubenzstein, et al., Phenotypic Characterization of Individuals with 30-40 CAG Repeats in the Huntington Disease (HD) Gene Reveals HD Cases with 36 Repeats and Apparently Normal Elderly Individuals with 36-39 Repeats," <u>Am. J. Hum. Genet.</u> , <u>59</u> , 16-22 (1996).
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JE		Takiyama et al., "The gene for Machado-Joseph disease maps to human chromosome 14q," <u>Nat. Genet.</u> , <u>4</u> , 300-304 (1993).
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JE		Trottier et al., "Polyglutamine expansions as a pathological epitope in Huntington's disease and four dominant cerebellar ataxias," <u>Letters to Nature</u> , <u>378</u> , 403-406 (1995).

EXAMINER Jasmine Enewold	Date Considered Jan 31, 2000
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